



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : D06M 23/00, 15/285, H01B 7/28		A1	(11) International Publication Number: WO 99/10591 (43) International Publication Date: 4 March 1999 (04.03.99)
<p>(21) International Application Number: PCT/EP98/05123</p> <p>(22) International Filing Date: 5 August 1998 (05.08.98)</p> <p>(30) Priority Data: 97202583.7 22 August 1997 (22.08.97) EP</p> <p>(71) Applicant (for all designated States except US): AKZO NOBEL N.V. [NL/NL]; Velperweg 76, NL-6824 BM Arnhem (NL).</p> <p>(72) Inventor; and</p> <p>(75) Inventor/Applicant (for US only): WILLEMSEN, Stephanus [NL/NL]; Methorstweg 11, NL-6991 TW Rheden (NL).</p> <p>(74) Agent: SCHALKWIJK, Pieter, Cornelis; Akzo Nobel N.V., Patent Dept. (Dept. APTA), P.O. Box 9300, NL-6800 SB Arnhem (NL).</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</p>	
<p>(54) Title: PROCESS FOR MANUFACTURE OF SUPERABSORBENT-COATED YARN</p> <p>(57) Abstract</p> <p>A process for the manufacture of a yarn provided with a superabsorbent material having a swelling value of at least 60 and having hydrophilic properties which is capable of absorbing and retaining quantities of water, characterized in that an aqueous solution comprising a water-soluble pre-superabsorbent material is applied onto the yarn, after which the yarn is dried and heated in order to cross-link or to polymerize the water-soluble pre-superabsorbent material to the superabsorbent material. The superabsorbent provides water blocking capability to the yarn, so that it is suitable for use as a strength member in water tight optical communication cables.</p>			

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

PROCESS FOR MANUFACTURE OF SUPERABSORBENT-COATED YARN

The invention relates to a process for the manufacture of yarn provided with
5 a superabsorbent material.

Said yarn can be utilized in cables, more particularly optical communications cables, in this way functioning simultaneously as reinforcing member and as water blocking agent.

10

A yarn provided with a superabsorbent material and the manufacture of such a yarn are known from European Patent Applications 482,703 and 351,100, US patent 5,635,569, and PCT application WO 93/18223. According to these publications yarns are impregnated with a
15 superabsorbent material in a water-oil emulsion. After being impregnated the treated yarn is dried, so that a film is formed in and around the yarn's interstices. The resulting product is used as reinforcing material for the manufacture of a communications cable with water blocking properties.

20 However, there are drawbacks to the above-mentioned methods of applying a superabsorbent material to a cable or yarn surface. A major disadvantage is the release of organic materials, in particular isohexadecane, during the process of drying the yarn. Such organic materials form an unacceptable environmental burden, and isohexadecane
25 must be eliminated by burning or by treatment in a waste gas unit. Such step requires expensive equipment and uses substantial amounts of energy. It is therefore an advantage to eliminate the use of water-oil emulsions, and to apply materials which are water soluble.

30 The drawback to impregnating an aramid yarn with a superabsorbent material dispersed in an aqueous system is that, due to the superabsorbent material's high viscosity enhancing action, uniform feeding of it is extremely

difficult, if not impossible. Further, on account of the limited concentration of superabsorbent material in the impregnating liquid only a small quantity of it can be applied to the yarn per treatment. Another drawback to this method is that the comparatively large amount of impregnating liquid which is 5 applied to the yarn with the superabsorbent material has to be removed again by evaporation.

The present invention obviates the aforesaid drawbacks. Moreover, the invention provides an aramid multifilament yarn of excellent water 10 absorbing capacity without using organic solvents or excessive amount of water.

The invention consists of a process for the manufacture of a yarn provided with a superabsorbent material having a swelling value of at least 60 and having hydrophilic properties which is capable of absorbing and retaining 15 quantities of water, characterized in that an aqueous solution comprising a water-soluble pre-superabsorbent material is applied onto the yarn, after which the yarn is dried and heated in order to cross-link or to polymerize the water-soluble pre-superabsorbent material to the superabsorbent material.

20

Thus no superabsorbent material as such is longer used but a water-soluble precursor which after heat treatment polymerizes or cross-links to superabsorbent material. This process may be followed by a step wherein a finish is applied onto the yarn. Alternatively, the aqueous solution already 25 comprises the finish.

The yarn obtained must have a swelling value greater than 60. The swelling value is a measure of the quantity of water absorbed by the yarn when it is contacted with water in the liquid form. The method of experimentally 30 determining the swelling value will be illustrated in greater detail below. It was found that the yarn according to the invention has swelling values comparable to the above-mentioned prior art yarns. Preferably, the swelling

value of the yarn according to the invention is at least 80, more particularly at least 100, and more particularly still at least 150.

The term multifilament yarn has the meaning it customarily has in the art,
5 i.e., the fibers of which it is made are endless filaments. Textile Terms and
Definitions (1988), pages 289-290 is referred to in this respect. Within the
framework of the invention the fibers used in making the yarns according to
the invention are filaments which may have any linear density common in
actual practice, and yarns may be made up of any desired number of
10 endless filaments. Generally, the filaments or the yarns composed thereof
will have a linear density of 0.01 to 20 000 dtex, while the endless filament
yarns will be composed of 1 to 20 000 filaments.

As suitable types of fibers may be mentioned fibers of organic as well as
inorganic origin. The fibers of organic origin may be either natural or
15 synthetic. Examples of natural fibers include cellulose fibers such as
cotton, linen, jute, etc., and fibers of animal origin such as wool, silk, etc.
Examples of synthetic organic fibers include fibers of regenerated cellulose,
rayon, polyesters, aliphatic and aromatic polyamides, acrylonitrile,
polyolefins, polyvinyl alcohol, polyvinyl chloride, polyphenylene sulfide,
20 elastomers, and carbon. Examples of inorganic fibers include fibers of
glass, metals, silica, quartz, etc., ceramic fibers, and mineral wool. In
addition, fibers made up of mixtures of said materials or copolymers thereof
or mixtures of said fibers may be employed. The aforementioned types of
fibers and other ones suitable for use in the process according to the
25 invention have been described in Kirk-Othmer, Encyclopedia of Chemical
Technology, 3rd Edition, Vol. 10 (1980), pp. 148-197.

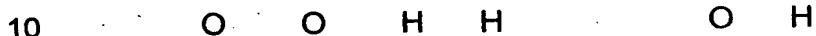
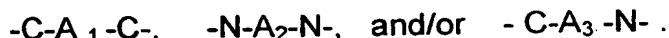
Preference is given to fibers composed of aramid, polyethylene
terephthalate, nylon-6, nylon-6,6, regenerated cellulose, or glass.

Also highly suitable as a substrate are fibers composed of two or more of
30 the aforementioned materials, e.g. bicomponent fibers. They may be of the
sheath-core or the side by side type, or of some other well-known type.

Other suitable types of fibers are satellite fibers and split fibers.

The fibers may be either solid or hollow. They may be round or flat or of any other desired cross-sectional shape, e.g. elliptical, triangular, star-shaped, kidney-shaped, etc.

5 Aramid yarns are most preferred. By aramid yarns is meant according to the invention yarns composed of polymers built up wholly or substantially from recurring units of the general formula:



Herein A_1 , A_2 , and A_3 represent different or the same divalent, one or more aromatic rings-containing rigid radicals which may also contain a heterocyclic ring, the chain-extending bonds of which radicals are in the position para to each other or are parallel and oppositely directed. 15 Examples of such radicals include 1,4-phenylene, 4,4'-biphenylene, 1,5-naphthylene, and 2,6-naphthylene. They may contain substituents or not, e.g. halogen atoms or alkyl groups. In addition to amide groups and the aforementioned aromatic radicals the chain molecules of the aromatic 20 polyamides may optionally contain up to 35 mole% of other groups, e.g. m-phenylene groups, non-rigid groups such as alkyl groups, or ether groups, urea groups or ester groups. Preferably, the aramid yarn is made up wholly or in part, more particularly for over 90 wt.%, of poly-(p-phenylene terephthalamide).

25 By a pre-superabsorbent material is meant according to the invention a material that after heating polymerizes or cross-links to a superabsorbent material having hydrophilic properties which is capable of absorbing and retaining a comparatively large quantity of water, optionally under pressure. 30 In order to simply apply the solution onto the yarn it is required that the pre-superabsorbent material dissolved in an aqueous solution has a viscosity smaller than $1000 \text{ mm}^2\text{sec}^{-1}$. Preferably, the viscosity is smaller than 500

$\text{mm}^2\text{sec}^{-1}$, and most preferably between 10 and 200 $\text{mm}^2\text{sec}^{-1}$. Particularly suitable for use as pre-absorbent materials are derivatives of polyacrylic acid. These include the water-soluble monomers, oligomers, and polymers derived from acrylamide, from acrylamide and sodium acrylate, and from 5 acrylamide and dialkylaminoethyl methacrylate. These compounds are selected from the groups of water-soluble non-ionic, anionic, and cationic monomers, oligomers, and polymers, respectively. Examples of pre-superabsorbents that can be used in the manufacture of the yarns according to the invention include: cross-linkable acrylic acid partially 10 neutralized into the sodium salt, polypotassium acrylate, co-oligo- or co-polymers of sodium acrylate and acrylamide, derivatives of co-polymers of acrylic acid and acrylamide, teroligo- or terpolymers of acrylamide and carboxyl groups and sulfo groups-containing monomers (sodium salt), polyacrylamide oligo- or polymers. Preferably, as pre-superabsorbent 15 material a derivative of polyacrylic acid or a derivative of a co-polymer of acrylamide and acrylic acid is used.

The term derivatives means that part of the carboxylic groups is in the salt form, preferably a potassium or sodium salt.

20 The yarns of the invention are made using a process wherein on the yarn surface there is provided a layer of a water solution comprising a pre-superabsorbent material, and that next the water is wholly or partially removed from the yarn by evaporation and thereafter or simultaneously heated to cross-link or polymerize the pre-superabsorbent material. 25 Optionally, a cross-linker or polymerization catalyst may be added to the aqueous solution of the pre-superabsorbent material.

30 The process to make the yarns according to the invention renders it possible to produce high quality aramid yarns having superabsorbent properties in an economical manner without using expensive equipment for after-burning organic solvents.

The amount of superabsorbent on the yarn is selected such that favorable water blocking properties are obtained when the yarn is used in cables. Favorable results are usually obtained if the yarn contains 0.3-10 wt.%, preferably 0.5-5 wt.%, and more particularly 0.6-2 wt.% of the 5 superabsorbent material.

The water solution having been applied, the yarn is dried. In this process the water is wholly or for the most part removed from the yarn by evaporation, so that a uniform layer of the superabsorbent material is left on 10 the yarn surface.

Drying takes place according to the conventional methods, in which use may be made of means such as hot drums, hot sheets, hot rollers, hot gases, tube ovens, steam boxes, infrared radiators, and the like. The drying temperature is 50 to 300°C. Heating is performed at 100 to 300°C. 15 Preferably, drying and heating is performed simultaneously at 100 to 250°C.

The process for obtaining the yarns according to the invention can be carried out in several ways.

20 In the fully continuous manner, which is linked directly to the yarn spinning process, the water solution containing the pre-superabsorbent can be applied to the washed yarn, after which the thus treated yarn is dried and heated. According to another embodiment, treatment of the yarn with the pre-superabsorbent present in a water solution, takes places in a separate 25 process not integrated with the spinning process.

The process for obtaining the yarns according to the invention is particularly suited to be used for combining, in one and the same process step, a heat treatment to cross-link or polymerize the pre-absorbent material, such as is 30 employed in the production of high-modulus aramid yarns.

In Kirk-Othmer, Encyclopedia of Chemical Technology, 3rd Ed., Vol. 3 (1978), pp. 213-242 a summary of aromatic polyamide preparation and spinning is provided. A pre-eminently suitable technique for wet spinning poly(p-phenylene terephthalamide) is described in US Patent Specification 5 4,320,081.

The aramid yarns according to the invention can have any linear density and any number of endless filaments common in actual practice. Generally, the yarn will have a linear density of 10 to 20 000 dtex and be composed of 10 10 to 20 000 filaments.

The superabsorbent material applied to the yarn according to the invention does not have a negative effect on the filament's principal mechanical properties.

15 The tenacity of the aramid yarns according to the invention is 1.0-3.5 N/tex or higher, preferably 1.5-2.5 N/tex. The elongation at break is 0.5-10%, preferably 1-8%. The initial modulus is 20-130 N/tex or higher, preferably 30-90 N/tex.

20 Like the corresponding yarn not treated with a superabsorbent, the yarn according to the invention has a relatively low water content after drying, even after it has been exposed to the air for a longer period of time. After the dried aramid yarn has been conditioned at 20°C and a relative humidity of 65% for 4 weeks, it has a water content not exceeding 10 wt.% , more 25 particularly not exceeding 7 wt.%.

The procedure to determine the swelling value of the yarn according to the invention is as follows.

About 10 g of the yarn to be examined are cut into non-intertwisted fibers of 30 some 12 cm in length.

The fiber sample is immersed completely, without stirring, in 600 ml of demineralized water of 20-22°C in an 800 ml beaker. For 60 seconds

(measured by stopwatch) the fiber sample remains immersed in the water in a condition of complete rest, i.e. without being stirred, shaken, vibrated, or subjected to some other form of movement. Immediately afterwards the overall contents of the beaker, i.e. fibers and water, are transferred to a bag 5 (dimensions: about 10 cm x 15 cm) made of polyester curtain netting (mesh size 1.5 mm x 1 mm). In this process the water for the most part runs out through the meshes of the curtain netting, while the fibers are left in the bag. Next, the bag and its contents are straight away transferred to a centrifuge and then centrifuged for 120 seconds (measured by stopwatch), 10 thus removing the still adhering water from the impregnated fiber sample. The centrifuge used is an AEG of the type SV 4528 (ex AEG Aktiengesellschaft, D-8500 Nuremberg), operates at a rate of 2800 revolutions per minute, and has a centrifugal drum with an inner diameter of about 24 cm. Immediately after it has been centrifuged the fiber sample is transferred 15 from the bag to a weighing box with a pair of tweezers and weighed to an accuracy of 0.0001 g (fiber weight: a grams). The fiber sample in the weighing box is thereupon dried to constant weight in an air oven at 105°C. Usually a drying time of 24 hours will suffice. After that the weight of the dried fiber sample in the weighing box is determined to an accuracy of 20 0.0001 g (fiber weight: b grams).

The swelling value of the yarn is calculated by means of the following formula:

$$25 \quad \text{swelling value} = \frac{(a-b) \cdot 100}{b}$$

Each determination is carried in duplicate and the results are averaged. Because of the properties mentioned herein before the yarn obtained 30 according to the invention is extremely well suited to be used as a reinforcing yarn of high water blocking capacity. Consequently, it is pre-eminently suitable for use in cables, more particularly optical communications cables.

The invention will be further illustrated with reference to the following examples.

Example I

5 A 19%-solution of sodium polyacrylate in water (Mirox W 60336, ex Stockhausen GmbH, Krefeld, Germany, viscosity $122 \text{ mm}^2\text{sec}^{-1}$ determined with Ubbelohde meter) was further dissolved in demineralized (demin) water. The solution was applied onto unfinished Twaron® yarn (1680 dtex/f 1000) with a dosage pump and an applicator. After drying in a steam box (length 10 m, 200-220°C, 15-60 sec) the swelling values were determined.

10 The content of Mirox on the yarn was determined to be 2 wt.%.

The results are depicted in Table I:

wt.% of Mirox solution	42.1	42.1	42.1	42.1	42.1	42.1	79.0	79.0	79.0
wt.% of demin water	57.9	57.9	57.9	57.9	57.9	57.9	21.0	21.0	21.0
concentration of Mirox in solution (%)	8.0	8.0	8.0	8.0	8.0	8.0	15.0	15.0	15.0
heating (sec)*	30	40	50	60	15	20	30	40	50
heating (°C)	200	200	200	200	220	220	200	200	200
yarn speed (m/min)	20	15	12	10	40	30	20	15	12
swelling value	210	139	78	60	192	81	178	107	66

* in steam box

15

At shorter heating times no cross-linking or polymerization occurs, and consequently swelling values are lower than 60.

Alternatively, the samples can be dried and heated in a tube oven (length 67 cm, 200°C), or on a hot-plate (length 42 cm, 150°C).

20

Example II

A 15%-solution of sodium acrylate/acrylamide copolymer in water (Produkt W 63194, ex Stockhausen GmbH, Krefeld, Germany, viscosity $191 \text{ mm}^2\text{sec}^{-1}$ determined with Ubbelohde meter) was further dissolved in demineralized (demin) water, and optionally Glyoxal (aq. glyoxal dialdehyde solution; ex Sigma-Aldrich Chemie bv, Zwijndrecht, The Netherlands), Sarpifan MKV (modified melamine resin solution in water; ex Stockhausen

10

GmbH, Krefeld, Germany), Breox 50A50 (ethoxylated and propoxylated butanol; ex INSPEC, Southampton, UK), and/or Leomin AN (ethoxylated phosphonate potassium salt; ex Clariant, Frankfurt, Germany) were added (see Table II). The solution was applied onto unfinished Twaron® yarn 5 (1680 dtex/f 1000) with a dosage pump and an applicator. After drying in a steam box (length 10 m) the swelling values were determined. The tables show the influence of the amount of finish on the yarn, and the effect of the residence time and the temperature.

10 Table II: compositions

	A	B	C	D	E	F	G	H
wt.% Produkt solution	75.0	50.0	25.0	12.5	75.0	50.0	50.0	50.0
Concentration of Produkt in solution (%)	11.25	7.5	3.75	1.875	11.25	7.5	7.5	7.5
wt.% Glyoxal (50%)	1.5	1.0	0.5	0.25			1.0	1.0
wt.% Sarpifan (50%)					1.5	1.0		
wt.% Breox							4.0	
wt.% Leomin								4.0
wt.% of demin water	23.5	49.0	74.5	87.25	23.5	49.0	45.0	45.0

Table III: swelling values as the result of residence time

composition 2.0%*	steam box			swelling value (%)
	m/min	sec	°C	
B	75	8	200	263
B	40	15	200	233
B	30	20	200	242
B	20	30	200	239
B	15	40	200	301
B	12	50	200	316
B	10	60	200	306

*amount of finish composition on yarn

Table IV: swelling values as the result of the amount of finish

composition (%)*	steam box			swelling value (%)
	m/min	sec	°C	
no finish	15	40	200	25
D (0.5)	15	40	200	74
C (1.0)	15	40	200	134
B (2.0)	15	40	200	304
A (3.0)	15	40	200	343
F (2.0)	15	40	200	63
E (3.0)	15	40	200	72

*amount of finish composition on yarn

5

Table V: swelling values as the result of the temperature

composition 2.0%*	steam box			swelling value (%)
	m/min	sec	°C	
B	15	40	160	165
B	15	40	198	198
B	15	40	220	91
B	15	40	240	68
F	15	40	160	214
F	15	40	198	94

*amount of finish composition on yarn

10 Table VI: swelling values as the result of the residence time

composition 2.0%*	steam box			swelling value (%)
	m/min	sec	°C	
F	75	8	200	219
F	40	15	200	85
F	20	30	200	71
F	10	60	200	61

*amount of finish composition on yarn

Table VII: swelling values as the result of the surfactant and residence time

composition 3.0%*	steam box			swelling value (%)
	m/min	sec	°C	
G	75	8	200	265
H	75	8	200	32
G	75	8	240	164
H	75	8	240	26
H	40	15	240	353

*amount of finish composition on yarn

CLAIMS

1. A process for the manufacture of a yarn provided with a superabsorbent material having a swelling value of at least 60 and having hydrophilic properties which is capable of absorbing and retaining quantities of water, characterized in that an aqueous solution comprising a water-soluble pre-superabsorbent material is applied onto the yarn, after which the yarn is dried and heated in order to cross-link or to polymerize the water-soluble pre-superabsorbent material to the superabsorbent material.
2. The process according to claim 1, followed by a step wherein a finish is applied onto the yarn.
- 15 3. The process according to claim 1 wherein the aqueous solution further comprises a finish.
4. The process according to any one of claims 1-3 wherein a multifilament aramid or glass yarn is used.
- 20 5. The process according to any one of claims 1-4 wherein as water-soluble pre-superabsorbent material a derivative of polyacrylic acid or a derivative of a co-polymer of acrylamide and acrylic acid is used.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 98/05123

A. CLASSIFICATION OF SUBJECT MATTER
IPC 6 D06M23/00 D06M15/285 H01B7/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 6 D06M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 351 100 A (AMERICAN TELEPHONE & TELEGRAPH) 17 January 1990 see column 6, line 8-18 ---	1-5
X	US 5 635 569 A (SACKMANN GUENTER ET AL) 3 June 1997 see column 1, line 38 - line 41; claims 1-3 ---	1-5
A	EP 0 544 435 A (AMERICAN TELEPHONE & TELEGRAPH) 2 June 1993 see column 4, line 53 - column 5, line 2 ---	1
A	EP 0 784 116 A (AKZO NOBEL NV) 16 July 1997 see the whole document ---	1 -/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "Z" document member of the same patent family

Date of the actual completion of the international search

4 January 1999

Date of mailing of the international search report

13/01/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl
Fax: (+31-70) 340-3016

Authorized officer

Tamme, H-M

INTERNATIONAL SEARCH REPORT

In ational Application No

PCT/EP 98/05123

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 779 389 A (AKZO NOBEL NV) 18 June 1997 see the whole document -----	1
A	EP 0 482 703 A (AKZO NOBEL NV) 29 April 1992 cited in the application see the whole document -----	1-5

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 98/05123

Patent document cited in search report	Publication date	Patent family member(s)			Publication date
EP 0351100	A 17-01-1990	US 4913517 A	CA 1323672 A	DK 339889 A	03-04-1990 26-10-1993 12-01-1990
		JP 2061909 A	JP 2618485 B	US 5389442 A	01-03-1990 11-06-1997 14-02-1995
US 5635569	A 03-06-1997	DE 4429318 A	DE 59501354 D	EP 0697416 A	22-02-1996 05-03-1998 21-02-1996
		JP 8245719 A			24-09-1996
EP 0544435	A 02-06-1993	US 5249248 A	AU 655687 B	AU 2833092 A	28-09-1993 05-01-1995 05-08-1993
		CA 2080930 A	CN 1073546 A, B	DE 69222921 D	28-05-1993 23-06-1993 04-12-1997
		DE 69222921 T	DE 544435 T	DK 19837 A	05-03-1998 02-02-1998 31-03-1996
		ES 2109320 T	ES 2662155 B	JP 6084412 A	16-01-1998 08-10-1997 25-03-1994
		JP 9206723 A	MX 9206723 A	MX 245288 A	31-05-1993 26-03-1996
EP 0784116	A 16-07-1997	AT 158623 T	DE 69314166 D	DE 69314166 T	15-10-1997 30-10-1997 12-03-1998
		DK 630434 T	WO 9318223 A	EP 0630434 A	11-05-1998 16-09-1993 28-12-1994
		ES 2107009 T	ES 7504463 T	ES 2107009 T	16-11-1997 18-05-1995 09-07-1996
EP 0779389	A 18-06-1997	US 5534304 A	AT 158622 T	CA 2054248 A	15-10-1997 27-04-1992
		DE 69127738 D	DE 69127738 T	DE 69127738 D	30-10-1997 12-03-1998 29-04-1992
		EP 0482703 A	ES 2106760 T	ES 2106760 T	16-11-1997 18-09-1992
		JP 4263670 A	US 5342686 A	JP 4263670 A	30-08-1994
		US 5264251 A	US 5264251 A	US 5264251 A	23-11-1993
EP 0482703	A 29-04-1992	AT 158622 T	CA 2054248 A	DE 69127738 D	15-10-1997 27-04-1992 30-10-1997
		DE 69127738 T	DE 69127738 T	EP 0779389 A	12-03-1998 18-06-1997
		EP 0779389 A	ES 2106760 T	ES 2106760 T	16-11-1997 18-09-1992
		JP 4263670 A	US 5342686 A	JP 4263670 A	30-08-1994
		US 5264251 A	US 5264251 A	US 5264251 A	23-11-1993